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## Amendments to the Claims

- nucleic acid analyte in a sample comprising: a first and second base region regions having at least one ribonucleotide modified to include a 2'-O alkyl substitution to the ribofurancesyl moiety; and a second base region, wherein the first and second base regions hybridize capable of hybridizing to each other under nucleic acid assay conditions to form a hybrid containing at least one ribonucleotide modified to include a 2'-O-alkyl substitution to the ribofurancesyl moiety, wherein the hybrid is more stable than a hybrid formed between unmodified forms of the first and second base regions, and wherein the oligonucleotide forms a hybrid with the nucleic acid analyte but not with a non-targeted nucleic acid under nucleic acid assay conditions, such that the nucleic acid analyte can be detected.
- 423. (Currently Amended) The oligonucleotide of claim 422, wherein that portion of the first base region capable of forming a hybrid with the second base region under nucleic acid assay conditions includes a cluster of at least about 4 ribonucleotides modified to include a 2'-O-alkyl substitution to the ribofuranosyl moiety.
- of the first base region capable of forming a hybrid with the second base region under nucleic acid assay conditions includes at least one nucleotide which is not a ribonucleotide modified to include a 2'-O-alkyl substitution to the ribofuranosyl moiety.
- 425. (Currently Amended) The oligonucleotide of claim 422, wherein each nucleotide of that portion of the first base region capable of forming a hybrid with the second base region under nucleic acid assay conditions is a ribonucleotide modified to include a 2'-O-alkyl substitution to the ribofuranosyl moiety.

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- 426. (Previously Added) The oligonucleotide of claim 422, wherein each nucleotide of the oligonucleotide is a ribonucleotide modified to include a 2'-O alkyl substitution to the ribofuranosyl moiety.
- 427. (Previously Added) The oligonucleotide of claim 422, wherein the oligonucleotide includes a conjugate molecule.
- 428. (Previously Added) The oligonucleotide of claim described to the oligonucleotide at a site located within the cluster of the first base region.
- 429. (Currently Amended) The oligonucleotide of claim 422, wherein the oligonucleotide is up to about between 10 and 100 bases in length.
- 430. (Previously Added) The oligonucleotide of claim 422, wherein the oligonucleotide includes a reporter group.
- 431. (Previously Added) The oligonucleotide of claim 430, wherein the reporter group comprises a fluorescent molecule.
- 432. (Previously Added) The oligonucleotide of claim 422, wherein the nucleic acid analyte comprises RNA.
- 433. (Previously Added) The oligonucleotide of claim 432, wherein the nucleic acid analyte comprises ribosomal RNA.

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- 434. (Previously Added) The oligonucleotide of claim 422, wherein the oligonucleotide is a hybridization assay probe which forms a detectable hybrid with the nucleic acid analyte.
- 435. (Previously Added) The oligonucleotide of claim 422, wherein the oligonucleotide is an amplification primer for use in an amplification procedure.
- 436. (Previously Added) The oligonucleotide of claim 435, wherein the amplification procedure is a polymerase chain reaction method of amplification.
- 437. (Previously Added) The oligonucleotide of claim 435, wherein the amplification procedure is a transcription-based method of amplification.
- 438. (Previously Added) The oligonucleotide of claim 422, wherein the oligonucleotide is a target capture oligonucleotide.
- 439. (Previously Added) The oligonucleotide of claim 438, wherein the target capture oligonucleotide is immobilized by a solid support.
- 440. (Previously Added) The oligonucleotide of claim 422, wherein the 2'-O-alkyl substitution to the ribofuranosyl moiety is a 2'-O-methyl substitution.
- 441. (Currently Amended) A method for determining the presence of a nucleic acid analyte in a sample, the method comprising the steps of:
- a) providing to the sample an the oligonucleotide of claim 422: comprising:

  i) a first base region having at least one ribonucleotide modified to include a 2'-O-alkyl substitution to the ribofuranosyl moiety; and

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hybridize to each other under nucleic acid assay conditions to form a hybrid more stable than a hybrid formed between unmodified forms of the first and second base regions, and wherein the oligonucleotide forms a hybrid with the nucleic acid analyte but not with a non-targeted nucleic acid in the sample under nucleic acid assay conditions, such that the nucleic acid analyte can be detected;

- b) incubating the sample under conditions such that the oligonucleotide hybridizes to the nucleic acid analyte, if present; and
- c) determining whether the oligonucleotide has hybridized to the nucleic acid analyte.
- first base region capable of forming a hybrid with the second base region under nucleic acid assay conditions includes a cluster of at least about 4 ribonucleotides modified to include a 2'-O-alkyl substitution to the ribofuranosyl moiety.
- first base region capable of forming a hybrid with the second base region under nucleic acid assay conditions includes at least one nucleotide which is not a ribonucleotide modified to include a 2'-O-alkyl substitution to the ribofuranosyl moiety.
- that portion of the first base region capable of forming a hybrid with the second base region under nucleic acid assay conditions is a ribonucleotide modified to include a 2'-O-alkyl substitution to the ribofuranosyl moiety.

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- 445. (Withdrawn) The method of claim 441, wherein each nucleotide of the oligonucleotide is a ribonucleotide modified to include a 2'-O-alkyl substitution to the ribofuranosyl moiety.
- 446. (Withdrawn) The method of claim 441, wherein the oligonucleotide includes a conjugate molecule.
- 447. (Withdrawn) The method of claim 442, wherein the oligonucleotide includes a conjugate molecule joined to the oligonucleotide at a site located within the cluster of the first base region.
- 448. (Currently Amended) The method of claim 441, wherein the oligonucleotide is up to about between 10 and 100 bases in length.
- 449. (Withdrawn) The method of claim 441, wherein the oligon ucleotide includes a reporter group.
- 450. (Withdrawn) The method of claim 449, wherein the reporter group comprises a fluorescent molecule.
- 451. (Withdrawn) The method of claim 441, wherein the nucleic acid analyte comprises RNA.
- 452. (Withdrawn) The method of claim 451, wherein the nucleic acid analyte comprises ribosomal RNA.

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- (Withdrawn) The method of claim 441, wherein the oligonucleotide is a 453. hybridization assay probe which forms a detectable hybrid with the nucleic acid analyte.
- (Withdrawn) The method of claim 441, wherein the oligonucleotide is an 454. amplification primer used in an amplification procedure.
- (Withdrawn) The method of claim 454, wherein the amplification procedure 455. is a polymerase chain reaction method of amplification.
- (Withdrawn) The method of claim 454, wherein the amplification procedure 456. is a transcription-based method of amplification.
- (Withdrawn) The method of claim 441, wherein the oligonucleotide is a target 457. capture oligonucleotide.
- (Withdrawn) The method of claim 457, wherein the target capture 458. oligonucleotide is immobilized by a solid support.
- (Withdrawn) The method of claim 441 further comprising the step of 459. quantifying the nucleic acid analyte determined to be present in the sample.
- (Withdrawn) The method of claim 454 further comprising the step of 460. quantifying the nucleic acid analyte determined to be present in the sample.
- (Currently Amended) The method of claim 441, wherein step c) is indicative of the presence or absence of an organism or one or more members of a group of organisms at least one mircoorganism or virus in the sample.

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- 462. (Withdrawn) The method of claim 441; further comprising the step of providing to the sample a nuclease inhibitor other than a polynucleotide modified to include a 2'-O-alkyl substitution to the ribofuranosyl moiety of a ribonucleotide.
- 463. (Withdrawn) The method of claim 441, wherein the 2'-O alkyl substitution to the ribofuranosyl moiety is a 2'-O-methyl substitution.
- 464. (New) The oligonucleotide of claim 432, wherein a target sequence contained within the nucleic acid analyte includes a double-stranded region.
- 465. (New) The method of claim 451, wherein a target sequence contained within the nucleic acid analyte includes a double-stranded region.